## **AMENDMENTS TO THE CLAIMS**

Docket No.: L7350.0010

- 1. (Currently amended) A method of polyubiqutinating a nucleophosmin comprising [of] reacting the nucleophosmin with BRCA1-BARD1.
- 2. (Currently amended) A method of stabilizing a nucleophosmin comprising of polyubiqutination of polyubiquitinating nucleophosmin.
- 3. (Original) The method of Claim 1 or 2, wherein polyubiquitination is carried out *in vitro* or *in vivo*.
- 4. (Currently amended) A method of inhibiting polyubiquitination of nucleophosmin comprising [of] phosphorylating BARD1 using CDK2-cyclin E or CDK2-cyclin A.
- 5. (Currently amended) A method of degrading and/or dissociating BRCA1-BARD1 comprising [of] phosphorylating BARD1 using CDK2-cyclin E and/or CDK2-cyclin A.
- 6. (Currently amended) A method of inactivating ubiquitin ligase activity of a BRCA1-BARD1 comprising [of] phosphorylating BARD1 using CDK2-cyclin E and/or CDK2-cyclin A.
- 7. (Currently amended) The method according to any one of Claims 4 to 6, wherein the phosphorylation sites of BARD1 are at least three sites selected from the group consisting of S148, S251, S288 and T299.
- 8. (Currently amended) The method according to any one of Claims 4 to 6, wherein the phosphorylation sites of BARD1 are S148, S288 and T299.
- 9. (Currently amended) The method according to any one of Claims 4 to 6, wherein the phosphorylation sites of BARD1 are S148, S251, S288 and T299.

10. (Currently amended) A method of transporting BRCA1 from a nucleus to cytoplasm wherein comprising co-expressing BRCA1 and CDK2-cyclin E and/or CDK2-cyclin A are co-expressed.

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